

Spontaneous Ejaculations Associated with Aripiprazole

Oğuzhan EĞİLMEZ, Mustafa ÇELİK, Aysun KALENDEROĞLU

Department of Psychiatry, Adıyaman University School of Medicine, Adıyaman, Turkey

ABSTRACT

Sexual side effects are common with antipsychotic use. Spontaneous ejaculations without sexual arousal have been previously described with several typical and atypical antipsychotics. We report the case of a man who had spontaneous ejaculations after stopping risperidone and starting 30 mg/day aripiprazole. Spontaneous ejaculations ceased 3 days after decreasing the aripiprazole dose to 15 mg/day. He denied sexual fantasies or increased sexual desire during the period in which he had spontaneous ejaculations. The partial agonistic effect of aripiprazole on D2 receptors may have augmented

the mesolimbic dopaminergic pathway, which was suppressed by risperidone, causing spontaneous ejaculations in this patient. Serotoninergic effects of aripiprazole should also be considered. This unusual side effect should be questioned, particularly in patients who recieve aripiprazole after D2-blocking antipsychotics; otherwise, this side effect may cause embarrassement and noncompliance.

Keywords: Aripiprazole, spontaneous ejaculation, risperidon

INTRODUCTION

Sexual side effects due to antipsychotic drugs are commonly observed (1). Impotence and erectile dysfunction are the most frequent sexual side effects, but spontaneous ejaculations without sexual arousal has also been described previously with zuclopentixol (2), trifluoperazin and thiotixen (3), risperidon (4), and ziprasidon (5). To the best of our knowledge, there is no report associating aripiprazole with spontaneous ejaculations.

CASE

Mr A was 24 years old and was married for 5 years. He was being followed at a psychiatry outpatient clinic for 3 years, with a diagnosis of paranoid schizophrenia. He was using 4 mg/day risperidon for 3 years. His delusions were controlled with this treatment. After his marriage, he complained of erectile dysfunction and loss of sexual interest, and we planned to change his treatment. Risperidon was stopped, and aripiprazole, which has a lower sexual side effect profile, was prescribed at a dose of 30 mg/day. He came to our clinic 2 weeks after the introduction of aripiprazole, complaining of spontaneous ejaculations without sexual arousal 3—4 times a day. He stated that ejaculations had started I week after the introduction of aripiprazole. He was consulted to the urology department, and no organic pathology was detected. He continued aripiprazole at the same dose for I more week and came to control again. He was still complaining of spontaneous ejaculations; therefore, the aripiprazole dosage was reduced to 15 mg/day. Spontaneous ejaculations stopped 3 days after dose reduction, and he denied loss of sexual interest or erectile dysfunction. In his follow-up, he used 15 mg/day aripiprazole for 6 months, and he did not report recurrence of spontaneous ejaculations. Remission in psychotic symptoms also sustained during follow-up. Written informed consent was taken from the patient in order to publish his data.

DISCUSSION

Antipsychotic drugs generally act to decrease libido by antagonising dopamine receptors and increasing prolactine levels (6,7). In contrast, dopamine agonism caused by certain drugs such as L-dopa, amphetamine, and pramipexole is associated with increased sexual desire and sexual arousal, which are reported by the patients as increased sexual fantasies, erections, and sexual activities (8). Brain dopamine systems between the hypothalamus and limbic system are thought to be the main sexual excitatory systems (9).

Aripiprazole is a unique antipsychotic drug because it causes a partial agonism at the dopamine D2 receptor (10). Risperidon decreases dopaminergic transmission and is therefore frequently associated with sexual side effects (11). Hence, aripiprazole may act to boost dopaminergic activity at the mesolimbic dopaminergic circuit, which was previously suppressed by risperidone. Aripiprazole also acts on serotoninergic receptors and has partial 5-HT $_{1A}$ agonist and 5HT $_{2A}$ antagonistic properties (12). Meston and Frohlich (13) reported that 5-HT $_{1A}$ agonism and 5HT $_{2A}$ antagonism promote sexuality. Lack of hyperprolactinemia in addition to these receptor effects may be the cause of spontaneous ejaculations in our patient, although further research is needed to clearly explain the sexual effects of aripiprazole.



Correspondence Address: Mustafa Çelik, Adıyaman Üniversitesi Tıp Fakültesi, Adıyaman, Türkiye E-mail: mustacelik@yahoo.com

Received: 30.07.2014 **Accepted:** 01.11.2014

Cheon et al. (14) reported two female patients who reported hypersexuality (demand for daily sexual intercourse, frequent use of online pornography, frequent masturbation, and unprovoked sexual urges toward strangers) after change in treatment regime from risperidone to aripiprazole. Vrignaud et al. (15) reported a male patient who had hypersexuality after aripiprazole treatment. Our patient had spontaneous erections and ejaculations after substituting aripiprazole for risperidone, but he denied increased sexual desire during these ejaculations.

Increased sexual desire and/or spontaneous sexual arousal should be questioned in schizophrenia patients taking aripiprazole, particularly if it is introduced after the cessation of D2 antagonists such as risperidone. If unrecognized, this unusual side effect may lead to embarrassment of the patient and noncompliance with treatment.

Conflict of Interest: No conflict of interest was declared by the authors.

Financial Disclosure: The authors declared that this study has received no financial support.

REFERENCES

- Aizenberg D, Zemishlany Z, Dorfman-Etrog P, Weizman A. Sexual dysfunction in male schizophrenic patients. | Clin Psychiatry 1995; 56:137-41.
- Gitlin MJ. Psychotropic medications and their effects on sexual function: diagnosis, biology, and treatment approaches. J Clin Psychiatry 1994; 55:406-13.
- Ischikawa J, Ishii H, Bonaccorso S, Fowler WL, OLaughlin IA, Meltzer HY. 5-HT2A and D2 receptor blockade increase cortical dopamine release via 5-HT1A receptor activation: a possible mechanism of atypical antipsychotic induced cortical dopamine release. J Neurochem 2001; 76:1521-1531. [CrossRef]

- Keck Jr P, Buffenstein A, Ferguson J, Feighner J, Jaffe W, Harrigan EP, Morrissey MR. Ziprasidone 40 and 120 mg/day in the acute exacerbation of schizophrenia and schizoaffective disorder: a 4-week placebo-controlled trial. Psychopharmacology (Berl) 1998; 140:173-184. [CrossRef]
- 5. Boora K, Chiappone K, Dubovsky S, Xu J. Ziprasidone-induced spontaneous orgasm. J Psychopharmacol 2010; 24:947-8. [CrossRef]
- Haddad PM, Wieck A. Antipsychotic-induced hyperprolactinaemia: mechanisms, clinical features and management. Drugs 2004; 64:229 I-23 I 4. [CrossRef]
- Knegtering H, van der Moolen AE, Castelein S, Kluiter H, van den Bosch RJ. What are the effects of antipsychotics on sexual dysfunctions and endocrine functioning? Psychoneuroendocrinology 2003; 28(Suppl 2):109-123. [CrossRef]
- 8. Sansone RA, Ferlan M. Pramipexole and compulsive masturbation. Psychiatry (Edgmont) 2007; 4:57-59.
- 9. Pfaus JG. Pathways of sexual desire. J Sex Med 2009; 6:1506-1533. [CrossRef]
- 10. Kessler RM. Aripiprazole: what is the role of dopamine D(2) receptor partial agonism? Am J Psychiatry 2007; 164:1310-1312. [CrossRef]
- 11. Wirshing DA, Pierre JM, Saunders CS, Wirshing WC. Sexual side effects of novel antipsychotic medications. Schizophr Res 2002; 56:25-30. [CrossRef]
- 12. Grunder G, Kungel M, Ebrecht M, Gorocs T, Modell S. Aripiprazole: pharmacodynamics of a dopamine partial agonist for the treatment of schizophrenia. Pharmacopsychiatry 2006; 39(Suppl I):S21-S25. [CrossRef]
- 13. Meston CM, Frohlich PF. The neurobiology of sexual function. Arch Gen Psychiatry 2000; 57:1012-1030. [CrossRef]
- 14. Cheon E, Koo BH, Seo SS, Lee JY. Two cases of hypersexuality probably associated with aripiprazole. Psychiatry Investig 2013; 10:200-202. [CrossRef]
- Vrignaud L, Aouille J, Mallaret M, Durrieu G, Jonville-Béra AP. Hypersexuality associated with aripiprazole: a new case and review of the literature. Therapie 2014; 69:525-527. [CrossRef]